# LUBRICANT EYE- carboxymethylcellulose sodium solution/drops The Kroger Co.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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# **Kroger Lubricant Eye Drops**

## Active ingredient

Carboxymethylcellulose sodium 0.5%

#### Purpose

Carboxymethylcellulose sodium.....Eye Lubricant

#### Uses

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun
- may be used as a protectant against further irritation

# □ Warnings

# For external use only.

**Do not use** if solution changes color or becomes cloudy.

# When using the product to avoid contamination

- do not touch tip of container to any surface.
- do not reuse.
- once opened, discard.
- do not touch unit-dose tip to eye.

# Stop use and ask a doctor if

- you experience eye pain.
- changes in vision occur.
- redness or irritation of the eye continues.
- redness or irritation of the eye worsens or persists for more than 72 hours.

**Keep out of reach of children.** If accidentally swallowed, get medical help or contact a Poison Control Center immediately.

### Directions

- To open, twist and pull tab to remove.
- Instill 1 or 2 drops in the affected eye(s) as needed and discard container.
- If used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions.

#### Other information

- Store at 15°-30°C (59°-86°F).
- Use only if single-use container is intact.
- Use before expiration date marked on container.
- RETAIN THIS CARTON FOR FUTURE REFERENCE

#### **Inactive ingredients**

Calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride, and sodium lactate. May contain sodium hydroxide and/or hydrochloric acid

### Questions or comments?

1-800-632-6900



### **LUBRICANT EYE**

carboxymethylcellulose sodium solution/ drops

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-373	
Route of Administration	ОРНТНАЬМІС			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 mL		

Inactive Ingredients	
Ingredient Name	Strength

CALCIUM CHLORIDE (UNII: M410 D6 VV5M)	
MAGNESIUM CHLO RIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660 YQ98110)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)	
HYDRO CHLO RIC ACID (UNII: QTT17582CB)	

1	Packaging				
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:30142-373- 33	30 in 1 BOX	06/01/2015		
1		0.5 mL in 1 VIAL, DISPENSING; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part349	06/01/2015		

# Labeler - The Kroger Co. (006999528)

# **Registrant** - Unimed Pharmaceuticals, Inc. (689852052)

Establishment			
Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	label(30142-373), manufacture(30142-373), pack(30142-373)

Revised: 12/2019 The Kroger Co.